

## DESCRIPTION

## COMPOSITION CONSISTING OF PLATELET-RICH PLASMA

## Technical Field

[0001]

The present invention relates to platelet-rich plasma to be used in the field of medical treatment and a method for the preparation thereof.

The present application claims the priority of Japanese Patent Application No. 2004-024815, which is incorporated herein by reference.

## Background Art

[0002]

Currently, medicine and engineering are making a great progress in the field of regeneration of lost tissue.

However, recently, the use of, for example, a blood product for medical use or a physiologically active substance produced from an animal (in particular, cow), which is contaminated with viruses ,prion and so on, are often cause infection, leading to a safety threatening events as a social problem. From this fact, a growing interest is focused on a safety concerning all medical treatments.

Further, for routine clinical applications, besides safety, reliability and convenience in operation are also

important.

[0003]

From such background, a fibrin formulation, which has an effect acknowledged as an enhancer for hemostasis and wound-healing after surgery, has been developed since early times to attract lots of attention on a therapeutic method for facilitating to heal a wound using self-blood components. For example, because an activated platelet secretes a substance which induces cell migration and differentiation at an early stage of wound healing, a plasma comprising the platelets concentrated from self-blood (platelet-rich plasma, hereinafter simply referred to as "PRP") is prepared before surgery and activated to apply onto an operation site, thereby to facilitate the therapy. Many clinicians have reported that this method has succeeded in facilitating wound healing.

[0004]

This method is a very safe method using self-blood and provides a reliable performance to some extent, but it has problems such as complexity to work, riskiness, laboriousness, necessity for skilled staffs, expensive devices to purchase and increasing cost for maintenance.

[0005]

Methods for preparing platelet-rich plasma, as well as methods and apparatus for separating blood into its components and the like have already been described in many

reports (patent document No. 1). Platelet-rich plasma for clinical testing can be obtained from a supernatant which is prepared by collecting blood from the median vein of forearm, adding each 4.5 mL of the whole blood into plastic tubes containing 1.5 mL of 3.1 w/v% sodium citrate, tumbling to mix, and then centrifuging by 500 g for 15 minutes at 22°C (non-patent document No. 1). Conventional centrifugation can precipitate erythrocytes, but needs a complicated work and can not be exempted from damaging hemocytes such as platelets and plasma proteins.

In the following, platelet-rich plasma obtained by a conventional method is referred to as "PRP" for convenience to differentiate from the platelet-rich plasma of the present invention.

[0006]

To obtain leukocyte-rich plasma (LRP), a method for adding both a rouleau formation aggregating agent and an enhancer to whole blood to separate erythrocyte cells from the whole blood is disclosed (patent document No. 2). Here, as the rouleau formation aggregating agent, for example, dextran, Hespan, Pentaspan and Ficoll are mentioned, and as the enhancer, oxalate, malonate, mannitol and sucrose are mentioned.

[0007]

Platelet-rich plasma used as an enhancer for hemostasis or wound-healing after surgery is desired to

comprise not only a platelet component but also much blood components including plasma proteins such as fibrinogen and leukocytes in the most same possible state as they are present in a living body. There is desired a platelet-rich plasma having so high an activity.

[0008]

Even though PRP is obtained by centrifugation or precipitation of hemocytes from blood, the PRP must contain not only platelets at least with their activity already enhanced but also a coagulation reaction inducer such as thrombin to use as a tissue and/or organ repair enhancer.

[patent document No. 1] JP H5-203639-A

[patent document No. 2] JP H8-510322

[non-patent document No. 1] "Outline of Clinical Laboratory Testing", 31st ed., pp. 400

Disclosure of the Invention

(Problems to be solved)

[0009]

An object of the present invention is to provide easily and inexpensively a platelet-rich plasma which does not always need a coagulation reaction inducer such as thrombin to add and has an effectively high activity to use as a tissue and/or organ repair enhancer.

(Means to Solve the Problems)

[0010]

The present inventors studied strenuously and, as a

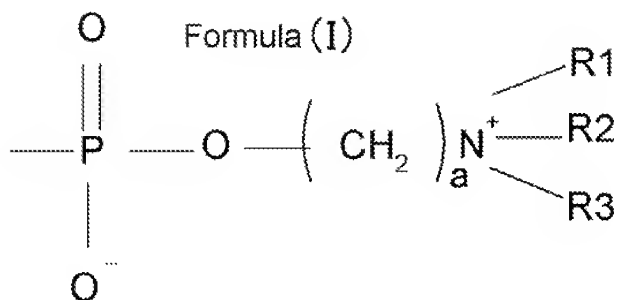
result, found that a polymer having a residue of an organic phosphate compound is added to a whole blood to aggregate and precipitate erythrocytes selectively and facilitatively, allowing provision of a platelet-rich plasma which comprises intact platelets similar to those in a living body and can be used as a tissue and/or organ repair enhancer free from a coagulation reaction inducer such as thrombin. The finding completes the present invention.

[0011]

Therefore, the present invention is composed of the following:

1. A method for preparing platelet-rich plasma, comprising adding a polymer having a residue of an organic phosphate compound to blood.
2. A method for preparing platelet-rich plasma, comprising the step of adding a polymer having a residue of an organic phosphate compound to blood to precipitate erythrocytes selectively.
3. The method for preparing platelet-rich plasma according to the preceding clause 1 or 2, comprising, as the polymer having a residue of an organic phosphate compound, a compound having in its side chain a group represented by the Formula (I) below:

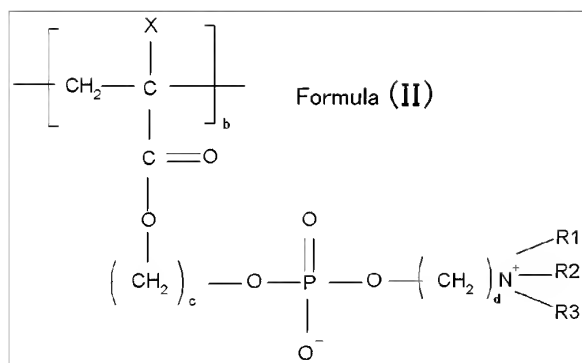
[Chem. 1]



(wherein R1, R2 and R3 are each independently a straight or branched chain alkyl group having a carbon number of 1 to 8, and a is an integer of 1 to 12).

4. The method for preparing platelet-rich plasma according to the preceding clause 3, comprising, as a polymer having a residue of an organic phosphate compound, a compound represented by the Formula (II) below:

[Chem. 2]

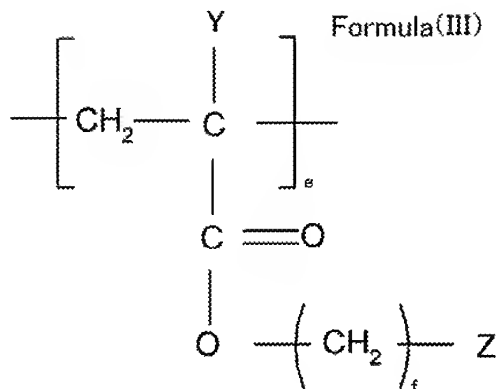


(wherein X is -H or -CH<sub>3</sub>, R1, R2 and R3 are each independently a straight or branched chain alkyl group having a carbon number of 1 to 8, b is an integer of 1 to 4,000, c is an integer of 1 to 6, and d is an integer of 1 to 12).

5. The method for preparing platelet-rich plasma

according to any one of the preceding clauses 1 to 4, further comprising, as a polymer having a residue of an organic phosphate compound, a compound represented by the Formula (III) below:

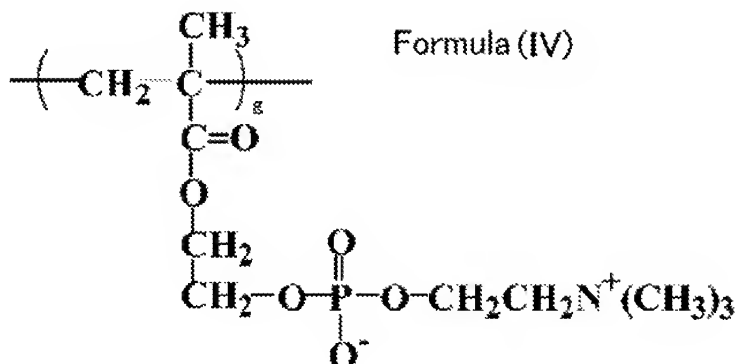
[Chem. 3]



(wherein Y is -H or -CH<sub>3</sub>, Z is -H, -OH, -COOH, -COONa, -COOK, -SO<sub>3</sub>H, -SO<sub>3</sub>Na, -SO<sub>3</sub>K, -NH<sub>2</sub>, -NHR<sub>1</sub>, -NR<sub>1</sub>R<sub>2</sub>, -N<sup>+</sup>R<sub>1</sub>R<sub>2</sub>R<sub>3</sub>OH<sup>-</sup>, -N<sup>+</sup>R<sub>1</sub>R<sub>2</sub>R<sub>3</sub>SO<sub>3</sub><sup>-</sup>, -N<sup>+</sup>R<sub>1</sub>R<sub>2</sub>R<sub>3</sub>NO<sub>3</sub><sup>-</sup>, -N<sup>+</sup>R<sub>1</sub>R<sub>2</sub>R<sub>3</sub>PO<sub>3</sub><sup>-</sup> [wherein R<sub>1</sub>, R<sub>2</sub> and R<sub>3</sub> are independently -(CH<sub>2</sub>)<sub>x</sub>H, wherein x is an integer of 1 to 5], -O-PO<sub>3</sub>Na, -O-PO<sub>3</sub>K, -O-PO<sub>3</sub>H, -OR<sub>4</sub>H, -COOR<sub>4</sub>H, -C=O-R<sub>4</sub>H, [wherein R<sub>4</sub> is aliphatic hydrocarbon, wherein the number of hydrocarbon is 0 to 20] or -O-R<sub>5</sub> [R<sub>5</sub> is an aromatic hydrocarbon group], e is an integer of 1 to 4000, and f is an integer of 0 to 20).

6. The method for preparing platelet-rich plasma according to the preceding clause 4 or 5, comprising, as a polymer having a residue of an organic phosphate compound, a compound represented by the Formula (IV) below:

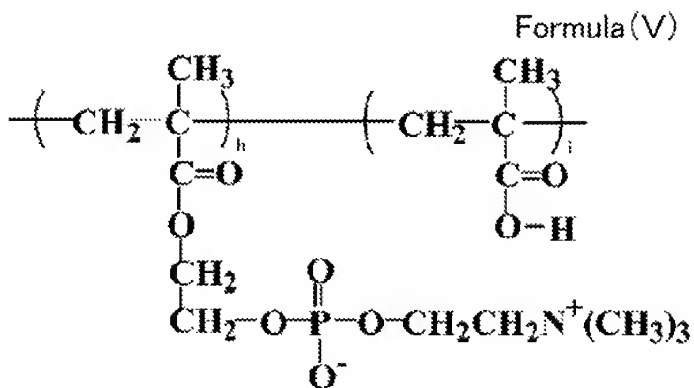
[Chem. 4]



(wherein g is an integer of 1 to 4,000).

7. The method for preparing the platelet-rich plasma according to any one of the preceding clauses 3 to 6, comprising, as a polymer having a residue of an organic phosphate compound, a compound represented by any of the Formulae (V) to (VII) below:

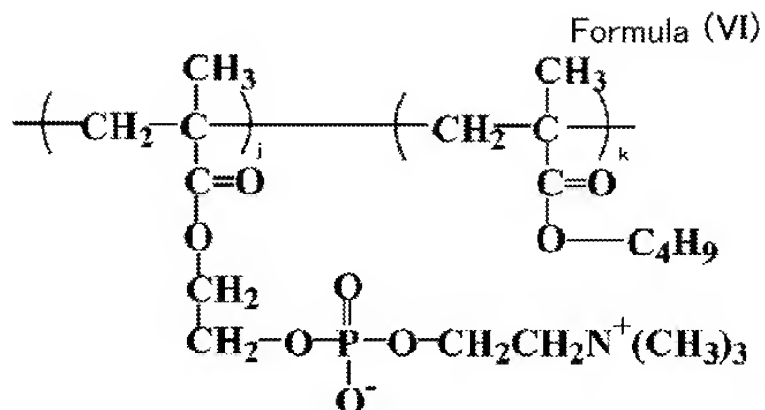
[Chem. 5]



(wherein h and i are each independently an integer of 1 to 4,000),



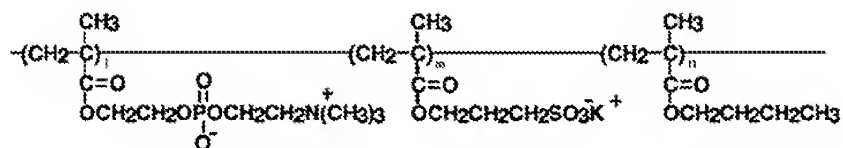
[Chem. 6]



(wherein j and k are each independently an integer of 1 to 4,000), and

[Chem. 7]

Formula (VII)



(wherein l, m and n are each independently an integer of 1 to 4,000).

8. Platelet-rich plasma obtained by the method for preparing according to any one of the preceding clauses 1 to 7.

9. A tissue and/or organ repair enhancer comprising the platelet-rich plasma according to the preceding clause 8.

10. A tissue and/or organ repair enhancer, augmentation

additive of bone surrounding dental implant, additive when implanting bone or artificial bone into a bone deficient part, wound healing enhancer, tissue healing enhancer after therapy or treatment for plastic and/or cosmetic purpose, skin disease therapeutic agent, skin ulcer therapeutic agent, nerve tissue repair agent and/or tissue repair agent after surgery comprising the platelet-rich plasma according to the preceding clause 8.

11. Any of the following therapeutic methods or treatment methods, comprising administering the platelet-rich plasma according to the preceding clause 8:

- 1) augmenting bone surrounding dental implant,
- 2) skin disease,
- 3) repairing tissue for plastic and/or cosmetic purpose,
- 4) repairing bone deficient part,
- 5) repairing nerve tissue, and
- 6) repairing tissue after surgery.

12. A polymer for preparing the platelet-rich plasma according to the preceding clause 8.

13. A reagent or a reagent kit for preparing platelet-rich plasma comprising the polymer according to the preceding clause 12.

14. A device for preparing the platelet-rich plasma according to the preceding clause 8.

(Effects of Invention)

[0012]

A polymer having a residue of an organic phosphate compound of the present invention is added to a blood to aggregate and precipitate erythrocytes more selectively than other hemocytes, allowing provision of platelet-rich plasma comprising a platelet which is little damaged and exists in its extremely intact state. The platelet-rich plasma, therefore, can be used as an effective tissue and/or organ repair enhancer. Further, the platelet-rich plasma obtained by the present method can be used with a reduced amount of a coagulation reaction inducer such as thrombin or to be free from the inducer as the case may be.

#### Brief Description of the Drawings

[0013]

Figure 1 shows the states of precipitated erythrocytes after the polymer was added to blood and allowed to stand for 20 minutes and 30 minutes. (Example 7)

Figure 2 shows platelet recovery rates of both platelet-rich plasma of the present invention and the PRP obtained by centrifugation of a conventional method. (Example 8)

Figure 3 shows amounts of fibrinogen contained in the platelet-rich plasma of the present invention and the PRP obtained by centrifugation of a conventional method. (Example 8)

Figure 4 shows activated-platelet rates (CD62P) in

the platelet-rich plasma of the present invention and the PRP obtained by centrifugation of a conventional method.

(Example 8)

#### Description of the Preferred Embodiment

[0014]

Platelets adhere and aggregate to damaged tissue to form thrombus, and play an extremely important role for hemostasis and further for storing and releasing substances which induce other cells to migrate and differentiate. Fibrin is produced by the action of thrombin on fibrinogen in plasma and involved in the final stage of blood coagulation. Further, fibrin is important as a scaffold for cells to infiltrate and differentiate in order to repair tissue. Further, it is known that leukocytes serve to prevent bacteria and harmful microorganisms from invading and have an immune function and a biocidal function, and in particular monocytes/macrophages among them play a key role in tissue repair.

[0015]

Platelets rapidly lose its function as time passes after stimulation or blood collection. Thereby, any other approach than centrifugation which little stimulates platelets and can separate quickly could not provide platelets holding functions to some extent. It is known that whole blood is allowed to stand to precipitate erythrocytes gradually, but that the rate is usually so

little that no other blood can be left for several hours or more to provide a platelet-containing plasma than a sick blood such as a blood in a high level inflammatory reaction.  
[0016]

The method for preparing platelet-rich plasma of the present invention can precipitate erythrocytes selectively and facilitatively from whole blood to give a supernatant which contains components essential for tissue repair not only platelets but also leukocytes and fibrinogen in their almost damage-free states with their respective activities held. Further, the platelet-rich plasma obtained by the present invention comprises platelets which are not almost activated to take the most possible intact state. The platelets in the most possible intact state have characters such as high hemostatic and coagulation effects and also maintain the high capacity to release a growth factor, so that platelet-rich plasma comprising such platelets has an excellent quality.

[0017]

By the present invention, erythrocytes can be selectively and facilitatively aggregated and precipitated. 80% or more, preferably 90% or more of erythrocytes contained in whole blood can be precipitated by aggregation within three hours, preferably within two hours, more preferably within an hour, and even more preferably within 30 minutes after starting the treatment, thereby to provide

platelet-rich plasma of at least 15% or more of total blood volume.

[0018]

Whole blood for use in the preparation of platelet-rich plasma of the present invention may be obtained according to the usual method for blood collection in human or animals other than human. Further, since the collected whole blood is left intact to coagulate, an anticoagulant is desired to add early after blood collection. As the anticoagulant, any may be used as long as it is commonly used and shows no toxicity to a living body, thus for example, those generally used such as sodium citrate, anticoagulant acid-citric-dextrose (ACD), EDTA, heparin, low molecular heparin, Futhan, hirudin and argatroban may be used.

[0019]

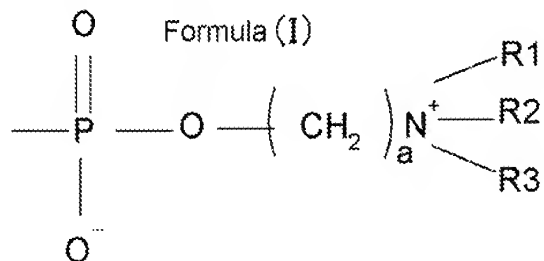
The amount of blood to be treated for preparing platelet-rich plasma of the present invention is not limited in particular and varied by usage, thus the adequate amount can be selected depending on the purpose or amount of use. The method of the present invention is applicable, for example, to self-blood as well as bloods collected from blood donors. As described above, the present invention is a method to conveniently obtain platelets with a high activity, thus it can also be used to prepare platelet formulations.

[0020]

(A polymer having a residue of an organic phosphate compound)

To selectively and facilitatively aggregate and precipitate erythrocytes from whole blood, a polymer having a residue of an organic phosphate compound can be added to whole blood. As the polymer having a residue of an organic phosphate compound, a compound comprising, as its side chain, a group represented by the Formula (I) below can be included in the constituent units:

[Chem. 8]



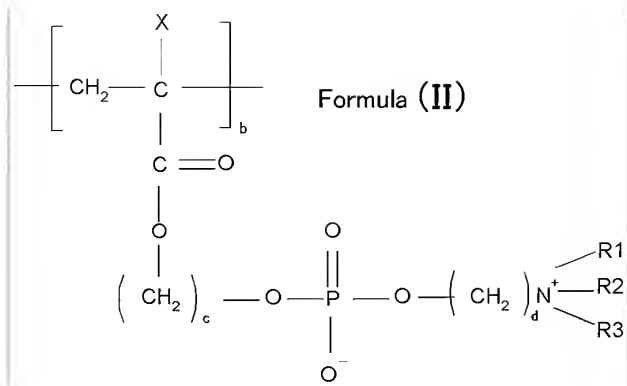
[0021]

(wherein R1, R2 and R3 are each independently a straight or branched chain alkyl group having a carbon number of 1 to 8, and a is an integer of 1 to 12).

[0022]

As the polymer having a residue of an organic phosphate compound, specifically a compound represented by the Formula (II) below can be included:

[Chem. 9]



[0023]

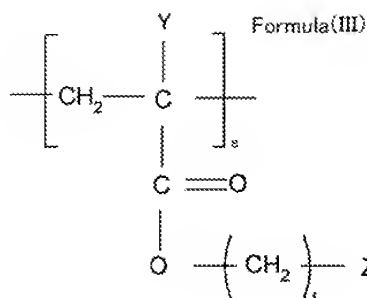
(wherein X is -H or -CH<sub>3</sub>, R1, R2 and R3 are each independently a straight or branched chain alkyl group having a carbon number of 1 to 8, b is an integer of 1 to 4000, c is an integer of 1 to 6, d is an integer of 1 to 12).

[0024]

Further, the polymer for use in the present invention may be a homopolymer comprising only one compound selected from Formula (II) as the constituent unit, or may be a heteropolymer comprising two or more compounds selected from Formula (II) as the constituent unit. Further, the polymer may be a heteropolymer comprising the other constituent units. Specifically, the polymer may include a methacrylate compound represented by Formula (III):

[Chem. 10]





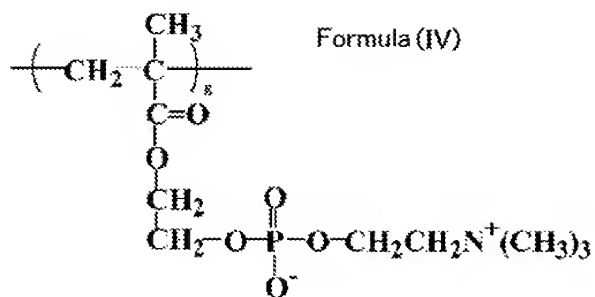
[0025]

(wherein Y is -H or -CH<sub>3</sub>, Z is -H, -OH, -COOH, -COONa, -COOK, -SO<sub>3</sub>H, -SO<sub>3</sub>Na, -SO<sub>3</sub>K, -NH<sub>2</sub>, -NHR<sub>1</sub>, -NR<sub>1</sub>R<sub>2</sub>, -N<sup>+</sup>R<sub>1</sub>R<sub>2</sub>R<sub>3</sub>OH<sup>-</sup>, -N<sup>+</sup>R<sub>1</sub>R<sub>2</sub>R<sub>3</sub>SO<sub>3</sub><sup>-</sup>, -N<sup>+</sup>R<sub>1</sub>R<sub>2</sub>R<sub>3</sub>NO<sub>3</sub><sup>-</sup>, -N<sup>+</sup>R<sub>1</sub>R<sub>2</sub>R<sub>3</sub>PO<sub>3</sub><sup>-</sup> [wherein R<sub>1</sub>, R<sub>2</sub> and R<sub>3</sub> are independently -(CH<sub>2</sub>)<sub>x</sub>H, wherein x is an integer of 1 to 5], -O-PO<sub>3</sub>Na, -O-PO<sub>3</sub>K, -O-PO<sub>3</sub>H, -OR<sub>4</sub>H, -COOR<sub>4</sub>H, -C=O-R<sub>4</sub>H, [wherein R<sub>4</sub> is aliphatic hydrocarbon, wherein the number of hydrocarbon is 0 to 20] or -O-R<sub>5</sub> [R<sub>5</sub> is an aromatic hydrocarbon group], e is an integer of 1 to 4000, and f is an integer of 0 to 20).

[0026]

As the polymer for use in the present invention, specifically a methacrylate polymer can be mentioned and, for example, 2-methacryloyl oxyethyl phosphorylcholine (MPC) represented by Formula (IV) below or derivatives thereof may be included:

[Chem. 11]



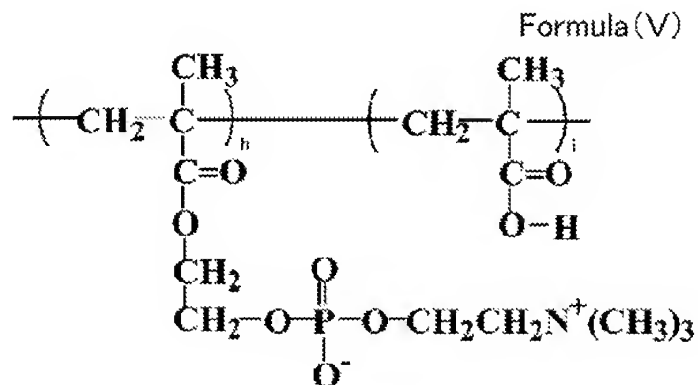
[0027]

(wherein g is an integer of 1 to 4000).

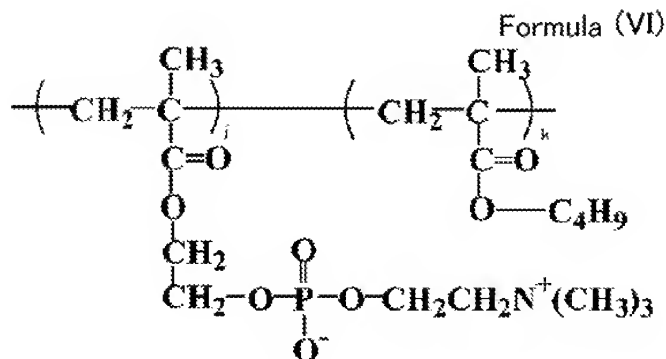
[0028]

As an example of heteropolymer for use in the present invention, compounds represented by the Formulae (V) and/or (VI) below can be included:

[Chem. 12]



[Chem. 13]



[0029]

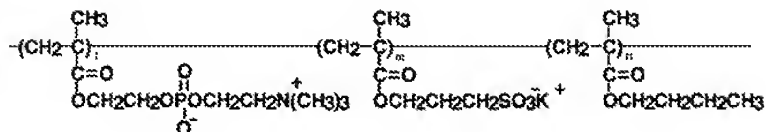
(wherein h, i, j and k are each independently an integer of 1 to 4000). For example, two kinds of constituent compounds represented in Formulae (V) and (VI) can have an appropriately selected ratio and, for example, 2-methacryloyl oxyethyl phosphorylcholine (MPC) and a methacrylate compound can have an appropriately selected constituent ratio such as 7:3, 8:2 and 9:1.

[0030]

Further, as an example of heteropolymer for use in the present invention, a compound represented by the Formula (VII) below can be included:

[Chem. 14]

Formula (VII)



[0031]

(wherein l, m and n are each independently an integer of 1 to 4000), and the represented three constituent compounds can have an appropriately selected ratio. 2-methacryloyl oxyethyl phosphorylcholine (MPC), Potassium 3-Methacryloyloxypropyl sulfonate (PMPS) and n-butyl methacrylate (BMA), which are contained in PMSB of Formula (VII), can have an appropriately selected ratio such as 25:20:55, 45:5:50 and 41:4:55. Meanwhile, for the purposes herein, PMSB25 means the polymer wherein the constituent ratio of MPC:PMPS:BMA is 25:20:55 and PMSB45 means the polymer wherein the constituent ratio of MPC:PMPS:BMA is 45:5:50.

[0032]

The average molecular weight of polymer applicable to the present invention is preferably in the range from 1,000 to 5,000,000, and more preferably from 1,000 to 1,000,000.

[0033]

(The method for preparing platelet-rich plasma)

To prepare the platelet-rich plasma of the present invention, into 1.5 mL of blood containing an anticoagulant, 0.0015 to 150 mg of, preferably 0.15 to 45 mg of, and more preferably 1.5 to 4.5 mg of the above described polymer having a residue of an organic phosphate compound can be added. They correspond to their respective concentrations

of approximately from 0.0001 to 10 w/v%, preferably from 0.01 to 3 w/v%, and more preferably from 0.1 to 0.3 w/v%. For preparing a larger amount of platelet-rich plasma, the blood volume should be increased and the polymer having a residue of an organic phosphate compound can be added to have the same rate as those described above. The above described polymer having a residue of an organic phosphate compound can be previously added in a container for holding collected blood, or directly in an injection syringe for blood collection.

[0034]

The polymer of an amount selected from the above volume range is added to blood containing an anticoagulant, and mixed gently to diffuse the polymer uniformly in blood, and then allowed to stand. Erythrocytes can be observed to precipitate selectively and facilitatively, thereby to give a supernatant, from which platelet-rich plasma can be obtained to comprise not only platelets but also a plasma component and a blood components such as leukocytes. Precipitation of erythrocytes can be observed in 3 hours at the longest, preferably within 2 hours, and as early as about 10 minutes after starting the treatment, so that the objective platelet-rich plasma of the present invention can be obtained within about 30 to 40 minutes.

Further, if erythrocytes are not sufficiently separated, or a shorter time of separation is needed,

conventional weak centrifugation can further be conducted to such an extent that no platelets may be aggregated. Specifically, centrifugation by 142 G for 5 minutes or less can be applied.

[0035]

(The use of platelet-rich plasma of the present invention)

The platelet-rich plasma prepared by the method of the present invention can be applied for any tissue or organ as a healing enhancer such as a wound healing agent, an augmentation additive of bone surrounding dental implant, an additive for implanting bone or artificial bone into a bone deficient part, a wound healing enhancer, a skin disease therapeutic agent, a skin ulcer therapeutic agent, a tissue healing enhancer after therapy or treatment for plastic or cosmetic purpose, a tissue repair agent after surgery, a tissue repair agent after surgery at an orthopedic operation site and a nerve tissue repair agent. Thus, with the platelet-rich plasma of the present invention, those disorders described above and damages in skins or tissues can be treated.

[0036]

The platelet-rich plasma of the present invention can be applied to human and mammals other than human. As an example of mammals, in particular, animals on the ground may be mentioned, and usually it can be applied to those kept as a pet such as dogs, cats and hamsters. Further,

it can be applied to animals employed in sports such as racehorses and bulls for fight.

[0037]

The platelet-rich plasma prepared by the method of the present invention from self-blood can be used for the purpose of therapy or treatment described above. Further, platelet-rich plasma, which is derived from the other blood than self-blood, can of course be used as long as the other blood is applicable in blood type.

[0038]

For use, specifically, any administration methods are applicable, and, for example, a required amount of the platelet-rich plasma of the present invention can be applied or injected to a wound site.

[0039]

(Preparative reagent, device and kit)

The platelet-rich plasma of the present invention is characterized in that, as described above, it can be prepared from self-blood conveniently and easily to be effective for therapy and the like. In order to achieve the convenient and easy preparation, a polymer having a residue of an organic phosphate compound for use in the preparation method of the present invention is stored as a reagent.

[0040]

Specifically, a reagent in which various polymers

explained in the section of "polymer having a residue of an organic phosphate compound" described above are filled in an appropriate container such as vial and ampoule, a plurality of these reagents, or a reagent kit comprising further a dissolution solution is also included in the present invention. Further, as a device for use in the preparation method of the present invention, specifically syringes and tubes for blood collection, and also sterilized containers or test tubes made of, for example, plastic, which are used to add a polymer having a residue of an organic phosphate compound to collected blood, may be mentioned. As the kit for preparing the platelet-rich plasma of the present invention, a kit which selects a reagent and a device selected from reagents and devices shown above to assemble may be mentioned. The present kit can be used to prepare the platelet-rich plasma of the present invention easily, for example, at the bedside where blood is collected.

[0041]

Further, syringes and containers for use in collecting blood, in which the polymer having a residue of an organic phosphate compound of the present invention has been filled in advance, are preferably used. A syringe or a container described above filled with the polymer having a residue of an organic phosphate compound is particularly preferably used, wherein the polymer has been dissolved in



sodium citrate or ACD which is broadly used as an anticoagulant.

[0042]

The present invention covers the platelet-rich plasma prepared by the method described above and a specific polymer in the present invention for use in preparing platelet-rich plasma.

Example

[0043]

In the following, the present invention will be explained in more detail with Examples and Experimental Examples, but the present invention will not be limited to them.

[0044]

Example 1

This Example aims to confirm effects of various monomers or polymers added to whole blood (peripheral blood) to precipitate erythrocytes.

[0045]

Seven samples were prepared by the following method. Anticoagulant acid-citrate-dextrose (hereinafter "ACD", from Shigma) was added to prepare a 3.13 w/v% ACD solution as an anticoagulant. 1.35 mL of blood was added to 0.15 mL of the ACD solution to prepare a whole blood sample (control) having a total of 1.5 mL.

[0046]

In this Example, for Sample 2, a heteropolymer Poly(MPC-co-BMA) (PMB (8:2)) (molecular weight of 100,000) comprising MPC and BMA in a ratio of 8:2 as described by formula VI was used, while for Sample 3, a homopolymer (molecular weight of 100,000) composed of only MPC as a constituent unit as described by formula IV was used. As a polyglutamic acid, the one from Shigma (molecular weight of 53,785) was used.

[0047]

Sample 1: Control

Sample 2: 3 mg of PMB (8:2) was dissolved in 0.15 mL of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 3: 3 mg of PMPC was dissolved in 0.15 mL of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 4: 0.1 mg of polyglutamic acid was dissolved in 0.15 mL of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 5: 0.5 mg of polyglutamic acid was dissolved in 0.15 mL of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 6: 0.7 mg of polyglutamic acid was dissolved in 0.15 mL of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 7: 3 mg of polyglutamic acid was dissolved in 0.15 mL of the ACD solution, and then mixed with 1.35 mL of blood.

[0048]

## 1) Precipitation of erythrocytes

The system of Sample 4 with polyglutamic acid added was almost equal to the systems of Samples 2 and 3 with the MPC-containing polymers added in Precipitation of erythrocytes after the samples were allowed to stand for 30 minutes.

[0049]

## 2) Platelet count

Leukocytes (WBC), erythrocytes (RBC), and platelets (PLT) contained in the supernatants after the samples were allowed to stand for 30 minutes were counted. Counting was conducted with an automatic hemocyte counter (Celltac $\alpha$  (MEC-6318)) supplied by Nihon Kohden co..

As a result, the supernatant obtained from Sample 3 showed a high value for leukocyte and platelet counts, like the supernatant obtained from Sample 4 (Table 1).

[Table 1]

|     | Sample1 | Sample2 | Sample3 | Sample4 | Sample5 | Sample6 | Sample7 |
|-----|---------|---------|---------|---------|---------|---------|---------|
| WBC | 65      | 124     | 140     | 140     | 100     | 97      | 88      |
| RBC | 397     | 6       | 13      | 5       | 5       | 3       | 4       |
| PLT | 17.3    | 26      | 41.6    | 37.4    | 33.6    | 33.9    | 31.9    |

[0050]

## Example 2

This Example aims to confirm the state of

precipitation of erythrocytes with or without an anticoagulant added at the time of adding PMSB represented by formula VII.

[0051]

Five samples were prepared by the following method. As in Example 1, 1.35 mL of blood was added to the ACD solution to prepare a whole blood sample (control) (Sample 1) having a total of 1.5 mL. Various weights of PMSB were dissolved in the ACD solution or the Hanks' solution (supplied by Invitrogen), and then added to blood to prepare Samples of 2 to 5. Here, PMSB25 in Samples 2 and 4 is a PMSB (molecular weight of 100,000) comprising MPC, PMPS and BMA described in formula VII in a ratio of 25:20:55, and PMSB45 in samples 3 and 5 is a PMSB (molecular weight of 100,000) comprising the compounds described in formula VII in a ratio of 45:5:50 likewise. Examples described below follow the same manner.

[0052]

Sample 1: Control

Sample 2: 3 mg of PMSB25 was dissolved in 0.15 ml of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 3: 3 mg of PMSB45 was dissolved in 0.15 ml of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 4: 3 mg of PMSB25 was dissolved in 0.15 ml of the Hanks' solution, and then mixed with 1.35 mL of blood.

Sample 5: 3 mg of PMSB45 was dissolved in 0.15 ml of

the Hanks' solution, and then mixed with 1.35 mL of blood.

[0053]

Each sample described above was allowed to stand for 30 minutes. In Samples 2 and 3, erythrocytes were observed to precipitate, thereby to give a supernatant, from which platelet-rich plasma could be obtained. On the other hand, in Samples 4 and 5 of systems without an anticoagulant contained, the samples were coagulated, and platelet-rich plasma could not be obtained.

[0054]

#### Example 3

This Example aims to examine effects of various PMSB added.

[0055]

The following 4 samples were prepared as in the method of Examples 1 and 2.

Sample 1: Control

Sample 2: 3 mg of polyglutamic acid was dissolved in 0.15 ml of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 3: 3 mg of PMSB25 was dissolved in 0.15 ml of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 4: 3 mg of PMSB45 was dissolved in 0.15 ml of the ACD solution, and then mixed with 1.35 mL of blood.

[0056]

Each sample described above was allowed to stand for

30 minutes. In samples 2 to 4, erythrocytes were observed to precipitate, thereby to give a supernatant, from which platelet-rich plasma was obtained. Leukocyte (WBC) and platelet (PLT) counts in the above described supernatant after standing, coagulation time and platelet activity (CD62P) were shown Table 2.

0.5 mL of supernatants of each sample, 0.05 mL of calcium chloride injection solution (as  $\text{CaCl}_2$ , 1.11 g/20 mL) in Pharmacopeia of Japan, and further 0.05 mL of self-blood with no anticoagulant were mixed in a 1.5 ml Eppendorf tube, and then the tube was slanted every 5 minutes to observe coagulation, and coagulation time was measured.

CD62P is a glycoprotein which is present in secreted granulosia in a platelet and has a molecular weight of 140kd, and once the platelet is activated, it moves to the surface of cell membrane of the platelet and expresses. Therefore, it serves as an indicator for activation of a platelet. CD62P was measured by flow cytometry supplied by BD Biosciences using an antibody.

[0057]

As a result, times needed for platelet-rich plasmas obtained from Samples 3 and 4 to coagulate were from 25 to 50 minutes, indicating that they had excellent coagulation ability in comparison to the platelet-rich plasma obtained from Sample 2. Further, the platelet-rich plasma obtained

from Sample 2 had a high CD62P value, indicating that the platelet had an enhanced activity, but the platelet-rich plasmas obtained from Samples 3 and 4 had low CD62P values (Table 2).

[Table 2]

#### Number of Blood Cells

|     | Sample 1 | Sample 2 | Sample 3 | Sample 4 |
|-----|----------|----------|----------|----------|
| WBC | 83       | 112      | 153      | 170      |
| PLT | 19.2     | 34.3     | 41.7     | 41.8     |

#### Coagulation Time

|      | Sample 1 | Sample 2                    | Sample 3 | Sample 4 |
|------|----------|-----------------------------|----------|----------|
| Time | ————     | Coagulated after 90 minutes | 50 min.  | 25 min.  |

#### CD62P (%Gated)

|        | Sample 1 | Sample 2 | Sample 3 | Sample 4 |
|--------|----------|----------|----------|----------|
| %Gated | * * *    | 95.73    | 4.58     | 4.42     |

[0058]

#### Example 4

This Example aims to examine effects of various PMSB added.

[0059]

The following four samples were prepared as in Example 3.

Sample 1: Control

Sample 2: 3 mg of polyglutamic acid was dissolved in 0.15 ml of the ACD solution, and then mixed with 1.35 mL

of blood.

Sample 3: 3 mg of PMSB25 was dissolved in 0.15 ml of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 4: 3 mg of PMSB45 was dissolved in 0.15 ml of the ACD solution, and then mixed with 1.35 mL of blood.

[0060]

Each sample described above was allowed to stand for 30 minutes. In Samples 2 to 4, erythrocytes were observed to precipitate to give a supernatant, from which the platelet-rich plasma was obtained. Leukocyte (WBC) and platelet (PLT) counts in the above described supernatant after standing and coagulation time were measured according to the method described in Examples 1 and 3.

As a result, times needed for platelet-rich plasmas obtained from Samples 3 and 4 to coagulate were from 10 to 20 minutes, indicating that they had excellent coagulation ability in comparison to platelet-rich plasma obtained from Sample 2 (Table 3).

[Table 3]

Number of Blood Cells

|     | Sample 1 | Sample 2 | Sample 3 | Sample 4 |
|-----|----------|----------|----------|----------|
| WBC | 89       | 138      | 161      | 160      |
| PLT | 22.1     | 41.6     | 45       | 46.4     |

Coagulation Time

|      | Sample 1 | Sample 2 | Sample 3 | Sample 4 |
|------|----------|----------|----------|----------|
| Time | ————     | 35min.   | 20min.   | 10min.   |



[0061]

Example 5

This Example aims to examine effects of various polymers added.

[0062]

The following five samples were prepared. 3.13 w/v% ACD solution of ACD added in water and the control of Sample 1 were prepared as in Example 1. For Samples 3 and 4, a homopolymer composed of only MPC as a constituent unit (molecular weight of 100,000) as described in formula IV were used, while for Sample 5, a heteropolymer comprising MPC and BMA in a ratio of 8:2 (PMB (8:2)) (molecular weight of 100,000) as described in formula VI was used. Various polymers were dissolved in water to get 5 w/v% as their polymer solutions.

[0063]

Sample 1: Control

Sample 2: 3mg of polyglutamic acid was dissolved in 0.15 ml of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 3: 3mg of polyglutamic acid was added to 0.06 mL of polymer solution (PMPC), supplied with 0.09 ml of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 4: 0.09 ml of the ACD solution was added to 0.06 mL of polymer solution (PMPC), and then mixed with 1.35 mL of blood.

Sample 5: 0.09 ml of the ACD solution was added to 0.06 mL of polymer solution (PMB (8:2)), and then mixed with 1.35 mL of blood.

[0064]

Leukocyte (WBC) and platelet (PLT) counts in the supernatants after the samples described above were allowed to stand for 30 minutes, coagulation times and activated-platelet rates (CD62P) were measured as in Examples 1 and 3.

As a result, it was confirmed that in the platelet-rich plasmas obtained from Samples 2 and 3 with polyglutamic acid contained, CD62P showed high values, while in the platelet-rich plasmas obtained from Samples 4 and 5, CD62P showed low values. Further, the platelet-rich plasmas obtained from Samples 4 and 5 coagulated within 10 minutes, while the platelet-rich plasmas obtained from Samples 2 and 3 with polyglutamic acid contained needed longer times for coagulation even the same amount of calcium chloride was added as in Samples 4 and 5. Further, the platelet-rich plasmas obtained from Samples 4 and 5, even if they were not supplied with a blood with no anticoagulant added, were supplied only with calcium chloride to coagulate within 10 minutes (Table 4).

[Table 4]

**Number of Blood Cells**

|            | Sample 1    | Sample 2    | Sample 3    | Sample 4    | Sample 5    |
|------------|-------------|-------------|-------------|-------------|-------------|
| <b>WBC</b> | <b>69</b>   | <b>103</b>  | <b>108</b>  | <b>142</b>  | <b>124</b>  |
| <b>PLT</b> | <b>21.3</b> | <b>45.3</b> | <b>43.8</b> | <b>51.6</b> | <b>42.9</b> |

**Coagulation Time**

|  | Sample 1 | Sample 2   | Sample 3   | Sample 4   | Sample 5   |
|--|----------|--|------------|--|--|
|  | * * *    | Not coagulated<br>after 35 min.<br>Afterwards<br>coagulated. | 10~35 min. | Within 10 min.<br>(Coagulated only<br>with CaCl <sub>2</sub> ) | Within 10 min.<br>(Coagulated only<br>with CaCl <sub>2</sub> ) |

**CD62P (%Gated)**

|                      | Sample 1     | Sample 2     | Sample 3     | Sample 4    | Sample 5    |
|----------------------|--------------|--------------|--------------|-------------|-------------|
| <b>After 30 min.</b> | <b>* * *</b> | <b>97.55</b> | <b>97.77</b> | <b>9.97</b> | <b>3.82</b> |

[0065]

**Example 6**

This Example aims to measure amount of fibrinogen contained in both platelet-rich plasmas obtained by the method of the present invention and PRP.

[0066]

The following three samples were prepared. Samples 1 and 2 were allowed to stand for 30 minutes according to the method of Example 1. For Sample 3, blood was added to 0.85 mL of aqueous solution with ACD3.8 w/v% mixed to prepare a solution having a total of 8.5 mL with the use of PRPkit (Kurasan KK), which was centrifuged by 3600 rpm for 15 minutes and then further by 2400 rpm for 10 minutes to give a supernatant.

[0067]

Sample 1: Control

Sample 2: 3mg of PMB (8:2) was dissolved in 0.15 mL

of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 3: Conventional method (centrifugation)

[0068]

In the supernatants obtained from the samples described above, the amounts of fibrinogens were measured. Measurement was conducted by the thrombin method using reagent from Boehringer Mannheim.

As a result, in Samples 1 and 2, high fibrinogen values were obtained. From this fact, it was confirmed that the platelet-rich plasmas obtained by the method of the present invention was neither denatured nor lost during operation, and thus had a large amount of fibrinogen (Table 5).

[Table 5]

|                       | Sample 1 | Sample 2 | Sample 3 |
|-----------------------|----------|----------|----------|
| Fibrinogen<br>(mg/dl) | 212      | 214      | 184      |

[0069]

#### Example 7

This Example aims to examine coagulation time when a heteropolymer represented by Formula (V) is added.

The following two samples were prepared. Sample 1 was allowed to stand for 20 and 30 minutes according to the method of Example 1 to give a supernatant. For Sample 2, 0.85 mL of 3.8 w/v% ACD in water was added to blood to give a solution having a total of 8.5 mL, which was then placed

into a 15 ml conical tube (BD Falcon tube) and centrifuged by 1000 rpm for 10 minutes to get a supernatant. The state of precipitation of erythrocytes in Sample 1 was shown in Figure 1.

[0070]

Sample 1: 3mg of both MPC and methacrylic acid copolymer (3:7) was dissolved in 0.15 mL of the ACD solution, and then mixed with 1.35 mL of blood.

Sample 2: Conventional method (centrifugation)

[0071]

Coagulation times were measured of the supernatants obtained from the samples described above. As in Example 3, 0.5 mL of the supernatant of each sample, 0.05 mL of calcium chloride injection solution (as  $\text{CaCl}_2$ , 1.11 g/20 mL) in Pharmacopeia of Japan, and further 0.05 mL of self-blood with no anticoagulant added were mixed in a 1.5 mL Eppendorf tube, and then the tube was slanted every 1 minute from 5 minutes after mixing to observe coagulation.

As a result, coagulation was observed in Sample 1 in 8 minutes, while Sample 2 obtained by a conventional method took 13 minutes to coagulate. From this fact, it was confirmed that the platelet-rich plasma obtained by the method of the present invention had excellent coagulation ability (Table 6).

[Table 6]

|                   | Sample 1 | Sample 2 |
|-------------------|----------|----------|
| Time to Coagulate | 8 min.   | 13 min.  |

[0072]

#### Example 8

This Example aims to examine platelet recovery rate, amount of fibrinogen and activated-platelet rate (CD62P) when PMB polymer is added.

The following two samples were prepared. Sample 1 was allowed to stand for 30 minutes according to the method of Example 5, yielding a supernatant. For sample 2, blood was added to 0.85 mL of aqueous solution with ACD3.8 w/v% mixed to prepare a solution having a total of 8.5 mL, which was placed in a 15 mL conical tube (BD Falcon tube) and centrifuged by 2400 rpm for 10 minutes, followed by collecting a supernatant with platelets contained and further centrifuging by 3600 rpm for 15 minutes to give a supernatant.

[0073]

Sample 1: 0.09 mL of ACD solution was added to 0.06 mL of polymer solution (PMB (8:2)), and then mixed with 1.35 mL of blood.

Sample 2: Conventional method (centrifugation)

[0074]

For supernatants obtained from the samples described above, platelet recovery rates, amount of fibrinogen and

CD62P were measured.

Platelet recovery rates were obtained according to the Formula below.

$$\{ (\text{amount of a supernatant} \times \text{platelet concentration in the supernatant}) / (\text{amount of blood collected} \times \text{platelet concentration at collecting blood}) \} \times 100 (\%)$$

CD62P was measured according to the method described in Example 3.

The amount of fibrinogen was measured according to the method described in Example 6.

[0075]

There were shown platelet recovery rate of supernatant obtained from each sample in Figure 2, amount of fibrinogen in Figure 3, and activated-platelet rate (CD62P) in Figure 4. As a result, in platelet recovery rate and amount of fibrinogen, the platelet-rich plasma obtained from Sample 1 was equivalent to that obtained from Sample 2 (conventional method). Further, in activated-platelet rate, the platelet-rich plasma obtained from Sample 1 was suppressed to have a lower value, and thus it was confirmed that this platelet-rich plasma had more excellent performance than the supernatant obtained in sample 2.

[Industrial Applicability]

[0076]

As explained above, platelet-rich plasma obtained by the preparation method of the present invention was

comparable to PRP obtained by conventional centrifugation in platelet recovery rate and amount of fibrinogen and kept a very high platelet activity. Therefore, platelet-rich plasma with excellent performance can be provided conveniently. In particular, platelet-rich plasma obtained by the method of the present invention using a self-blood component as the source may be employed in the field of medical treatment as a tissue and/or organ repair enhancer, in particular, an augmentation additive of bone surrounding dental implant, an additive for implanting bone or artificial bone into a bone deficient part, a wound healing enhancer, a tissue healing enhancer after therapy or treatment for plastic and/or cosmetic purpose, a skin disease therapeutic agent, a skin ulcer therapeutic agent and a nerve tissue repair agent and/or a tissue repair agent after surgery.